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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/520,801	08/10/2005	Zhikang Peng	82402-8502	9960
Michael R Williams Ade & Company 1700 360 Main Street Winnipeg Manitoba, R3C 3Z3 CANADA			EXAMINER	
			MERTZ, PREMA MARIA	
			ART UNIT	PAPER NUMBER
			1646	
	·		MAIL DATE	DELIVERY MODE
			08/10/2007	PAPER

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

	Application No.	Applicant(s)			
	10/520,801	PENG ET AL.			
Office Action Summary	Examiner	Art Unit			
	Prema M. Mertz	1646			
The MAILING DATE of this communication app Period for Reply	ears on the cover sheet with the c	orrespondence ad	ldress		
A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 1 MONTH(S) OR THIRTY (30) DAYS, WHICHEVER IS LONGER, FROM THE MAILING DATE OF THIS COMMUNICATION.  - Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.  - If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.  - Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).					
Status					
Responsive to communication(s) filed on  2a)    This action is <b>FINAL</b> .    2b)    This  3)    Since this application is in condition for allowar closed in accordance with the practice under E	action is non-final. nce except for formal matters, pro		e merits is		
Disposition of Claims					
4) ⊠ Claim(s) 1-42 is/are pending in the application. 4a) Of the above claim(s) is/are withdraw 5) □ Claim(s) is/are allowed. 6) □ Claim(s) is/are rejected. 7) □ Claim(s) is/are objected to. 8) ⊠ Claim(s) 1-42 are subject to restriction and/or expressions.	vn from consideration.				
Application Papers					
9) The specification is objected to by the Examiner 10) The drawing(s) filed on is/are: a) access applicant may not request that any objection to the Replacement drawing sheet(s) including the correction of the oath or declaration is objected to by the Examiner  11) The oath or declaration is objected to by the Examiner  12. **The Specification**  13. **The Specification**  14. **The Specification**  15. **The Specification**  16. **The Specification**  16. **The Specification**  17. **The Specification**  18. **The Specification**  19. **The Specification**  19. **The Specification**  11. **The Specification**  11. **The Specification**  11. **The Specification**  12. **The Specification**  13. **The Specification**  14. **The Specification**  15. **The Specification**  16. **The Specification**  16. **The Specification**  17. **The Specification**  17. **The Specification**  18. **The Specification**  19. **The Specification**  19. **The Specification**  19. **The Specification**  19. **The Specification**  10. **The Specification**  11. **The Specification**  11. **The Specification**  11. **The Specification**  12. **The Specification**  13. **The Specification**  13. **The Specification**  14. **The Specification**  15. **The Specification**  16. **The Specification**  16. **The Specification**  17. **The Specification**  18. **The Specification**  19. **The Specification**  10. **The Specification**  10. **The Specification**  11. **The Specification**  12. **The Specification**  13. **The Specification**  14. **The Specification**  15. **The Specification**  16. **The Speci	epted or b) objected to by the Eddrawing(s) be held in abeyance. See ion is required if the drawing(s) is obj	e 37 CFR 1.85(a). ected to. See 37 C			
Priority under 35 U.S.C. § 119					
<ul> <li>12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).</li> <li>a) All b) Some * c) None of:</li> <li>1. Certified copies of the priority documents have been received.</li> <li>2. Certified copies of the priority documents have been received in Application No.</li> <li>3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).</li> <li>* See the attached detailed Office action for a list of the certified copies not received.</li> </ul>					
Attachment(s)  1) Notice of References Cited (PTO-892)  2) Notice of Draftsperson's Patent Drawing Review (PTO-948)	4)  Interview Summary Paper No(s)/Mail Da 5)  Notice of Informal P	ate			
Information Disclosure Statement(s) (PTO/SB/08)     Paper No(s)/Mail Date	6) Other:	атол приовног			

#### **DETAILED ACTION**

#### Election/Restriction

1. This application is a 371 of PCT/CA04/00610. For applications filed under 371, PCT rules for lack of unity apply.

2. Restriction is required under 35 U.S.C. 121 and 372.

This application contains inventions or groups of inventions, which are not so linked as to form a single inventive concept under PCT Rule 13.1.

In accordance with 37 CFR 1.499, applicant is required, in reply to this action, to elect a single invention to which the claims must be restricted.

Groups I-33. Claims 1-11, 40, drawn to a reagent comprising a cytokine or chemokine-derived peptide or a cytokine-receptor-derived peptide or chemokine-receptor derived-peptide; a carrier protein; and an adjuvant.

Groups 34-66. Claims 12-23, 41, drawn to a method of inducing an immune response by administering a reagent comprising a cytokine or chemokine-derived peptide or a cytokine-receptor-derived peptide or chemokine-receptor derived-peptide; a carrier protein; and an adjuvant.

Groups 67-99. Claims 24-35, 42, drawn to a method of treating or ameliorating asthma by administering a reagent comprising a cytokine or chemokine-derived peptide or a cytokine-receptor-derived peptide or chemokine-receptor derived-peptide; a carrier protein; and an adjuvant.

Groups 100-132. Claims 24-35, 42, drawn to a method of preventing asthma by administering a reagent comprising a cytokine or chemokine-derived peptide or a cytokine-

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receptor-derived peptide or chemokine-receptor derived-peptide; a carrier protein; and an adjuvant.

Groups 133-165. Claims 36-39, drawn to an expression system comprising a nucleic acid encoding a reagent comprising a cytokine or chemokine-derived peptide or a cytokine-receptor-derived peptide or chemokine-receptor derived-peptide; a carrier protein; and an adjuvant.

NOTE: Should any one of the Groups from 1-165 be elected, Applicants are required to select one polypeptide (one amino acid sequence) as set forth in claim 36. Once one polypeptide sequence is selected all other sequences will be withdrawn from consideration.

NOTE: Independent claim 1 and claim 2 reads on polypeptides that are not related in structure and function, and therefore these claims are considered to comprise improper Markush groups. Claim 1 is not a proper linking claim because it, in fact, comprises multitudes of polypeptide sequences.

Applicants must choose a single polypeptide sequence for examination. This is not a species election, but an election of a single invention.

If Applicants believe that their sequences are so overlapping as to be obvious variants of each other, Applicants may choose a single sequence for search, this sequence being a representative sequence of all sequences or a designated subset of the sequences, as Applicants may choose. If Applicants present a single sequence to represent all sequences claimed, it will be understood that if this sequence or any sequence is found, the remaining sequences will be considered to be obvious variants of the found sequence.

The inventions listed as Groups I-165 do not relate to a single general inventive concept under PCT Rule 13.1 because under PCT Rule 13.2 they lack the same or corresponding special technical feature for the following reasons:

The PCT rules define a special technical feature as a feature, which defines a contribution over the prior art. The first claimed invention fails to recite such a feature, since US patent 6,358,509 (Ramanathan et al, (2002)), teaches a IL-4 fragment SEQ ID NO:I (26 amino acids) which is 100% identical in 15 amino acid overlap to present SEQ ID NO:I (column 6, line 25 column 7, line 13; column 5, line 65 - column 6, line 7; Table 1, columns 11-12). Therefore, the IL-4 peptide of the reference meets the limitations of the reagent of Group I. Since the first claimed invention lacks a special technical feature, the other claimed inventions cannot share a special technical feature with the first claimed invention. The inventions of Groups 133-165 are patentably distinct from the products of Groups I-33 because the products of Groups 1-33 can be synthesized by materially different methods, such as by chemical synthesis. The inventions of Groups 1-33 are patentably distinct from the products of Groups 133-165 because the products of Groups 133-165 can be used in methods that are materially different from the methods in which the inventions of Groups 1-33 are used, such as in gene therapy. The methods of Groups 34-132 are patentably distinct from each other because each recites method steps not required by the other, each method uses different starting materials and the search of all methods in one patent application would result in an undue search burden.

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3. Applicant is advised that the response to this requirement to be complete must include an election of the invention to be examined even though the requirement be traversed (37 CFR 1.143).

Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 C.F.R. 1.48(b) if one or more of the currently named inventors is no longer an inventor of at least one claim remaining in the application. Any amendment of inventorship must be accompanied by a diligently-filed petition under 37 C.F.R. 1.48(b) and by the fee required under 37 C.F.R. 1.17(h).

## Rejoinder under In re Ochiai, In re Brouwer

4. The examiner has required restriction between product and process claims. Where applicant elects claims directed to the product, and a product claim is subsequently found allowable, withdrawn process claims that depend from or otherwise include all the limitations of the allowable product claim will be rejoined in accordance with the provisions of MPEP § 821.04. Process claims that depend from or otherwise include all the limitations of the patentable product will be entered as a matter of right if the amendment is presented prior to final rejection or allowance, whichever is earlier.

Amendments submitted after final rejection are governed by 37 CFR 1.116; amendments submitted after allowance are governed by 37 CFR 1.312.

In the event of rejoinder, the requirement for restriction between the product claims and the rejoined process claims will be withdrawn, and the rejoined process claims will be fully examined for patentability in accordance with 37 CFR 1.104. Thus, to be allowable, the rejoined claims must meet all criteria for patentability including the requirements of 35 U.S.C. 101, 102,

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103, and 112. Until an elected product claim is found allowable, an otherwise proper restriction requirement between product claims and process claims may be maintained. Withdrawn process claims that are not commensurate in scope with an allowed product claim will not be rejoined. See "Guidance on Treatment of Product and Process Claims in light of *In re Ochiai, In re Brouwer* and 35 U.S.C. § 103(b)," 1184 O.G. 86 (March 26, 1996). Additionally, in order to retain the right to rejoinder in accordance with the above policy, Applicant is advised that the process claims should be amended during prosecution either to maintain dependency on the product claims or to otherwise include the limitations of the product claims. **Failure to do so may result in a loss of the right to rejoinder.** 

Further, note that the prohibition against double patenting rejections of 35 U.S.C. 121 does not apply where the restriction requirement is withdrawn by the examiner before the patent issues. See MPEP § 804.01.

### **Advisory Information**

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Prema Mertz whose telephone number is (571) 272-0876. The examiner can normally be reached on Monday-Friday from 7:00AM to 3:30PM (Eastern time).

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Gary Nickol, can be reached on (571) 272-0835.

Official papers filed by fax should be directed to (571) 273-8300. Faxed draft or informal communications with the examiner should be directed to (571) 273-0876.

Information regarding the status of an application may be obtained from the Patent application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see <a href="http://pair-direct.uspto.gov">http://pair-direct.uspto.gov</a>. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

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Prema Mertz Ph.D., J.D. Primary Examiner Art Unit 1646 August 3, 2007